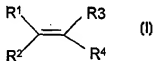




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(21) International Application Number: PCT/GB98/01702 (22) International Filing Date: 11 June 1998 (11.06.98) (30) Priority Data: <div style="display: flex; justify-content: space-between; margin-top: 5px;"> <div>9712338.4 9720078.6</div> <div>14 June 1997 (14.06.97) 23 September 1997 (23.09.97)</div> <div>GB GB</div> </div> (71) Applicant (for all designated States except US): THE SECRETARY OF STATE FOR DEFENCE [GB/GB]; Defence Evaluation & Research Agency, Ively Road, Farnborough, Hampshire GU14 0LX (GB). (72) Inventors; and (75) Inventors/Applicants (for US only): BADYAL, Jas, Pal, Singh [GB/GB]; University of Durham, Science Laboratories, South Road, Durham DH1 3LE (GB). COULSON, Stephen, Richard [GB/GB]; University of Durham, Science Laboratories, South Road, Durham DH1 3LE (GB). WILLIS, Colin, Robert [GB/GB]; CBDE, Porton Down, Salisbury, Wiltshire SP4 0JQ (GB). BREWER, Stuart, Anson [GB/GB]; CBDE, Porton Down, Salisbury, Wiltshire SP4 0JQ (GB). (74) Agent: SKELTON, S., R.; D/PR, Formalities Section (Procurement Executive), Poplar 2, MOD Abbey Wood # 19, Bristol BS34 8JH (GB).		(81) Designated States: AU, CA, CN, GB, JP, NZ, SG, SI, US, European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE). Published <i>With international search report. Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</i>
(54) Title: SURFACE COATINGS		
(57) Abstract <p>A method of coating a surface with a polymer layer, which method comprises exposing said surface to a plasma comprising a monomeric unsaturated organic compound which comprises a chain of carbon atoms, which are optionally substituted by halogen; provided that where the compound is a perhalogenated alkene, it has a chain of at least 5 carbon atoms; so as to form an oil or water repellent coating on said substrate. Suitable compounds for use in the methods are compounds of formula (I) where R¹, R² and R³ are independently selected from hydrogen, alkyl, haloalkyl or aryl optionally substituted by halo; provided that at least one of R¹, R² or R³ is hydrogen, and R⁴ is a group X-R⁵ where R⁵ is an alkyl or haloalkyl group and X is a bond; a group of formula -C(O)O(CH₂)_nY- where n is an integer of from 1 to 10 and Y is a bond or a sulphonamide group; or a group -(O)_pR⁶(O)_q(CH₂)_t- where R⁶ is aryl optionally substituted by halo, p is 0 or 1, q is 0 or 1 and t is 0 or an integer of from 1 to 10, provided that where q is 1, t is other than 0. The method is particularly useful in the production of oil- and/or water repellent fabrics.</p>		



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Surface Coatings

The present invention relates to the coating of surfaces, in particular to the production of oil- and water- repellent surfaces, as well as to coated articles obtained thereby.

Oil- and water- repellent treatments for a wide variety of surfaces are in widespread use. For example, it may be desirable to impart such properties to solid surfaces, such as metal, glass, ceramics, paper, polymers etc. in order to improve preservation properties, or to prevent or inhibit soiling.

A particular substrate which requires such coatings are fabrics, in particular for outdoor clothing applications, sportswear, leisurewear and in military applications. Their treatments generally require the incorporation of a fluoropolymer into or more particularly, fixed onto the surface of the clothing fabric. The degree of oil and water repellency is a function of the number and length of fluorocarbon groups or moieties that can be fitted into the available space. The greater the concentration of such moieties, the greater the repellency of the finish.

In addition however, the polymeric compounds must be able to form durable bonds with the substrate. Oil- and water- repellent textile treatments are generally based on fluoropolymers that are applied to fabric in the form of an aqueous emulsion. The fabric remains breathable and permeable to air since the treatment simply coats the fibres with a very thin, liquid-repellent film. In order to make these finishes durable, they are sometimes co-applied with cross-linking resins that bind the fluoropolymer treatment to fibres. Whilst good levels of durability towards

laundering and dry-cleaning can be achieved in this way, the cross-linking resins can seriously damage cellulosic fibres and reduce the mechanical strength of the material.

Chemical methods for producing oil- and water-repellent
5 textiles are disclosed for example in WO 97/13024 and British patent No 1,102,903 or M. Lewin et al., 'Handbook of Fibre Science and Technology' Marcel and Dekker Inc., New York, (1984) Vol 2, Part B Chapter 2.

10 Plasma deposition techniques have been quite widely used for the deposition of polymeric coatings onto a range of surfaces. This technique is recognised as being a clean, dry technique that generates little waste compared to conventional wet chemical methods. Using this method,
15 plasmas are generated from small organic molecules, which are subjected to an ionising electrical field under low pressure conditions. When this is done in the presence of a substrate, the ions, radicals and excited molecules of the compound in the plasma polymerise in the gas phase and react
20 with a growing polymer film on the substrate. Conventional polymer synthesis tends to produce structures containing repeat units which bear a strong resemblance to the monomer species, whereas a polymer network generated using a plasma can be extremely complex.

25 The success or otherwise of plasma polymerisation depends upon a number of factors, including the nature of the organic compound. Reactive oxygen containing compounds such as maleic anhydride, has previously been subjected to plasma
30 polymerisation (Chem. Mater. Vol. 8, 1, 1996).

US Patent No 5,328,576 describes the treatment of fabric or paper surfaces to impart liquid repellent properties by

subjecting the surfaces to a pre-treatment with an oxygen plasma, followed by plasma polymerisation of methane.

However, plasma polymerisation of the desirable oil and water repellent fluorocarbons have proved more difficult to achieve. It has been reported that cyclic fluorocarbons undergo plasma polymerisation more readily than their acyclic counterparts (H. Yasuda et al., J. Polym. Sci., Polym. Chem. Ed. 1977, 15, 2411). The plasma polymerization of trifluoromethyl-substituted perfluorocyclohexane monomers has been reported (A. M. Hynes et al., Macromolecules, 1996, 29, 18-21).

A process in which textiles are subjected to plasma discharge in the presence of an inert gas and subsequently exposed to an F-containing acrylic monomer is described in SU-1158-634. A similar process for the deposition of a fluoroalkyl acrylate resists on a solid substrate is described in European Patent Application No. 0049884.

Japanese application no. 816773 describes the plasma polymerisation of compounds including fluorosubstituted acrylates. In that process, a mixture of the fluorosubstituted acrylate compounds and an inert gas are subjected to a glow discharge.

The applicants have found an improved method of producing polymer and particular halopolymer coatings which are water and/or oil repellent on surfaces.

According to the present invention there is provided a method of coating a surface with a polymer layer, which method comprises exposing said surface to a plasma comprising a monomeric unsaturated organic compound which

comprises an optionally substituted hydrocarbon group, wherein the optional substituents are halogen; provided that where the compound is a straight chain perhalogenated alkene, it includes at least 5 carbon atoms; so as to form
5 an oil or water repellent coating on said substrate.

Unsaturated organic compounds are those which contain at least one double bond which is capable of reacting to form a polymeric compound. The compounds used in the method of
10 the invention suitably include at least one optionally substituted hydrocarbon chain. Suitable chains, which may be straight or branched, have from 3 to 20 carbon atoms, more suitably from 6 to 12 carbon atoms

15 Monomeric compounds used in the method may include the double bond within a chain and so comprise alkenyl compounds. Alternatively, the compounds may comprise an alkyl chain, optionally substituted by halogen, as a substituent which is attached to an unsaturated moiety
20 either directly or by way of a functional group, such as a ester or sulphonamide group.

As used therein the term "halo" or "halogen" refers to fluorine, chlorine, bromine and iodine. Particularly
25 preferred halo groups are fluoro. The term hydrocarbon includes to alkyl, alkenyl or aryl groups. The term "aryl" refers to aromatic cyclic groups such as phenyl or naphthyl, in particular phenyl. The term "alkyl" refers to straight or branched chains of carbon atoms, suitably of up to 20
30 carbon atoms in length. The term "alkenyl" refers to straight or branched unsaturated chains suitably having from 2 to 20 carbon atoms.

Monomeric compounds where the chains comprise unsubstituted alkyl or alkenyl groups are suitable for producing coatings which are water repellent. By substituting at least some of the hydrogen atoms in these chains with at least some halogen atoms, oil repellency may also be conferred by the coating.

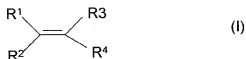
Thus in a preferred aspect, the monomeric compounds include haloalkyl moieties or comprise haloalkenyls. Therefore, preferably the plasma used in the method of the invention will comprise a monomeric unsaturated haloalkyl containing organic compound.

Suitable plasmas for use in the method of the invention include non-equilibrium plasmas such as those generated by radiofrequencies (Rf), microwaves or direct current (DC). They may operate at atmospheric or sub-atmospheric pressures as are known in the art.

The plasma may comprise the monomeric compound alone, in the absence of other gases or in mixture with for example an inert gas. Plasmas consisting of monomeric compound alone may be achieved as illustrated hereinafter, by first evacuating the reactor vessel as far as possible, and then purging the reactor vessel with the organic compound for a period sufficient to ensure that the vessel is substantially free of other gases.

Particularly suitable monomeric organic compounds are those of formula (I)

6



where R^1 , R^2 and R^3 are independently selected from hydrogen, alkyl, haloalkyl or aryl optionally substituted by halo; and
 5 R^4 is a group $\text{X}-\text{R}^5$ where R^5 is an alkyl or haloalkyl group and X is a bond; a group of formula $-\text{C}(\text{O})\text{O}(\text{CH}_2)_n\text{Y}-$ where n is an integer of from 1 to 10 and Y is a bond or a sulphonamide group; or a group $-(\text{O})_p\text{R}^6(\text{O})_q(\text{CH}_2)_t-$ where R^6 is aryl optionally substituted by halo, p is 0 or 1, q is 0 or
 10 1 and t is 0 or an integer of from 1 to 10, provided that where q is 1, t is other than 0.

Suitable haloalkyl groups for R^1 , R^2 , R^3 and R^5 are fluoroalkyl groups. The alkyl chains may be straight or
 15 branched and may include cyclic moieties.

For R^5 , the alkyl chains suitably comprise 2 or more carbon atoms, suitably from 2-20 carbon atoms and preferably from 6 to 12 carbon atoms.

20

For R^1 , R^2 and R^3 , alkyl chains are generally preferred to have from 1 to 6 carbon atoms.

Preferably R^5 is a haloalkyl, and more preferably a
 25 perhaloalkyl group, particularly a perfluoroalkyl group of formula $\text{C}_m\text{F}_{2m+1}$ where m is an integer of 1 or more, suitably from 1-20, and preferably from 6-12 such as 8 or 10.

Suitable alkyl groups for R^1 , R^2 and R^3 have from 1 to 6
 30 carbon atoms.

Preferably however, at least one of R^1 , R^2 and R^3 is hydrogen and preferably R^1 , R^2 , R^3 are all hydrogen.

Where X is a group $-C(O)O(CH_2)_nY-$, n is an integer which provides a suitable spacer group. In particular, n is from 1 to 5, preferably about 2.

Suitable sulphonamide groups for Y include those of formula $-N(R^7)SO_2$ where R^7 is hydrogen or alkyl such as C_{1-6} alkyl, in particular methyl or ethyl.

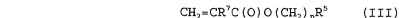
In a preferred embodiment, the compound of formula (I) is a compound of formula (II)



where R^5 is as defined above in relation to formula (I).

In compounds of formula (II), X in formula (I) is a bond.

In an alternative preferred embodiment, the compound of formula (I) is an acrylate of formula (III)



where n and R^5 as defined above in relation to formula (I) and R^7 is hydrogen or C_{1-6} alkyl, such as methyl.

Using these compounds, coatings with water hydrophobicity values of up to 10 and oleophobicity values of up to 8 have been achieved as illustrated hereinafter.

Other compounds of formula (I) are styrene derivatives as are well known in the polymer art.

All compounds of formula (I) are either known compounds or they can be prepared from known compounds using conventional methods.

5

The surface coated in accordance with the invention may be of any solid substrate, such as fabric, metal, glass, ceramics, paper or polymers. In particular, the surface comprises a fabric substrate such as a cellulosic fabric, to which oil- and/or water-repellency is to be applied. Alternatatively, the fabric may be a synthetic fabric such as an acrylic/nylon fabric.

The fabric may be untreated or it may have been subjected to earlier treatments. For example, it has been found that treatment in accordance with the invention can enhance the water repellency and confer a good oil-repellent finish onto fabric which already has a silicone finish which is water repellent only.

20

Precise conditions under which the plasma polymerization takes place in an effective manner will vary depending upon factors such as the nature of the polymer, the substrate etc. and will be determined using routine methods and/or the techniques illustrated hereinafter. In general however, polymerisation is suitably effected using vapours of compounds of formula (I) at pressures of from 0.01 to 10 mbar, suitably at about 0.2mbar.

25

A glow discharge is then ignited by applying a high frequency voltage, for example at 13.56MHz.

The applied fields are suitably of average power of up to 50W. Suitable conditions include pulsed or continuous

fields, but are preferably pulsed fields. The pulses are applied in a sequence which yields very low average powers, for example of less than 10W and preferably of less than 1W. Examples of such sequences are those in which the power is
5 on for 20 μ s and off for from 10000 μ s to 20000 μ s.

The fields are suitably applied for a period sufficient to give the desired coating. In general, this will be from 30 seconds to 20 minutes, preferably from 2 to 15 minutes,
10 depending upon the nature of the compound of formula (I) and the substrate etc.

Plasma polymerisation of compounds of formula (I), particularly at low average powers has been found to result
15 in the deposition of highly fluorinated coatings which exhibit super-hydrophobicity. In addition, a high level of structural retention of the compound of formula (I) occurs in the coating layer, which may be attributed to the direct polymerisation of the alkene monomer for instance a
20 fluoroalkene monomer via its highly susceptible double bond.

It has been noted, particularly in the case of the polymerisation of compounds of formula (III) above, that low power pulsed plasma polymerisation produces well-adhered
25 coatings which exhibit excellent water and oil repellency. The greater level of structural retention in the case of pulsed plasma polymerisation can be attributed to free radical polymerisation occurring during the duty cycle off-time and less fragmentation during the on-time.

30

In a particularly preferred embodiment of the invention, a surface is exposing a surface to a plasma comprising a compound of formula (III) as defined above, wherein the

plasma being created by a pulsed voltage also as described above.

Suitably the compound of formula (I) includes a
5 perfluoroalkylated tail or moiety, the process of the invention may have oleophobic as well as hydrophobic surface properties.

Thus the invention further provides a hydrophobic or
10 oleophobic substrate which comprises a substrate comprising a coating of a alkyl polymer and particularly a haloalkyl polymer which has been applied by the method described above. In particular, the substrates are fabrics but they may be solid materials such as biomedical devices.

15 The invention will now be particularly described by way of example with reference to the accompanying diagrammatic drawings in which:

20 Figure 1 shows a diagram of the apparatus used to effect plasma deposition;

Figure 2 is a graph showing the characteristics of
continuous wave plasma polymerisation of 1H, 1H, 2H-
25 pefluoro-1-dodecene;

Figure 3 is a graph showing the characteristics of pulsed
plasma polymerisation of 1H, 1H, 2H-pefluoro-1-dodecene at
50W P_p , $T_{on}=20\mu s$ and $T_{off}=10000\mu s$ for 5 minutes; and

30 Figure 4 is a graph showing the characteristics of (a) continuous and (b) pulsed plasma polymerisation of 1H, 1H, 2H, 2H-heptafluorodecyl acrylate.

Example 1Plasma Polymerisation of Alkene

1H, 1H, 2H-perfluoro-1-dodecene ($C_{10}F_{21}CH=CH_2$) (Fluorochem
5 F06003, 97% purity) was placed into a monomer tube (1) (Fig.
1) and further purified using freeze-thaw cycles. A series
of plasma polymerisation experiments were carried out in an
inductively coupled cylindrical plasma reactor vessel (2) of
5cm diameter, 470cm³ volume, base pressure of 7×10^{-3} mbar, and
10 with a leak rate of better than 2×10^{-3} cm³min⁻¹. The reactor
vessel (2) was connected by way of a "viton" O-ring (3), a
gas inlet (4) and a needle valve (5) to the monomer tube
(1).

15 A thermocouple pressure gauge (6) was connected by way of a
Young's tap (7) to the reactor vessel (2). A further
Young's tap (8) connected with an air supply and a third (9)
lead to an E2M2 two stage Edwards rotary pump (not shown) by
way of a liquid nitrogen cold trap (10). All connections
20 were grease free.

An L-C matching unit (11) and a power meter (12) was used to
couple the output of a 13.56 Mhz R.F. generator (13), which
was connected to a power supply (14), to copper coils (15)
25 surrounding the reactor vessel (2). This arrangement
ensured that the standing wave ratio (SWR) of the
transmitted power to partially ionised gas in the reactor
vessel (2) could be minimised. For pulsed plasma
deposition, a pulsed signal generator (16) was used to
30 trigger the R.F power supply, and a cathode ray oscilloscope
(17) was used to monitor the pulse width and amplitude. The
average power <P> delivered to the system during pulsing is
given by the following formula:

$$\langle P \rangle = P_{cw} \{ T_{on} / (T_{on} + T_{off}) \}$$

where $T_{on} / (T_{on} + T_{off})$ is defined as the duty cycle and P_{cw} is the average continuous wave power.

5

In order to carry out polymerization/deposition reactions the reactor vessel (2) was cleaned by soaking overnight in a chlorox bleach bath, then scrubbing with detergent and finally rinsing with isopropyl alcohol followed by oven
10 drying. The reactor vessel (2) was then incorporated into the assembly as shown in Figure 1 and further cleaned with a 50W air plasma for 30 minutes. Next the reactor (2) vessel was vented to air and the substrate to be coated (19), in this case a glass slide, was placed in the centre of the
15 chamber defined by the reactor vessel (2) on a glass plate (18). The chamber was then evacuated back down to base pressure (7.2×10^{-3} mbar).

Perfluoroalkene vapour was then introduced into the reaction
20 chamber at a constant pressure of ~0.2mbar and allowed to purge the plasma reactor, followed by ignition of the glow discharge. Typically 2-15 minutes deposition time was found to be sufficient to give complete coverage of the substrate. After this, the R.F generator was switched off and the
25 perfluoroalkene vapour allowed to continue to pass over the substrate for a further 5 minutes before evacuating the reactor back down to base pressure, and finally venting up to atmospheric pressure.

30 The deposited plasma polymer coatings were characterised immediately after deposition by X-ray photoelectron spectroscopy (XPS). Complete plasma polymer coverage was confirmed by the absence of any Si (2p) XPS signals showing through from the underlying glass substrate.

- A control experiment, where the fluoroalkene vapour was allowed to pass over the substrate for 15 minutes and then pumped down to base pressure was found to show the presence of a large Si (2p) XPS signal from the substrate. Hence the coatings obtained during plasma polymerisation are not just due to absorption of the fluoroalkene monomer onto the substrate.
- The experiments were carried out with average powers in the range of from 0.3 to 50W. The results of the XPS spectrum of a 0.3W continuous wave plasma polymer deposition onto a glass slide for 13 minutes is shown in Figure 2.
- It can be seen that in this instance, CF_2 and CF_3 groups are the prominent environments in the C(1s) XPS envelope:-

CF_2 (291.2eV) 61%

CF_3 (293.3eV) 12%

20

The remaining carbon environments comprised partially fluorinated carbon centres and a small amount of hydrocarbon (C_xH_y). The experimental and theoretically expected (taken from the monomer) values are given in Table 1

25

Table 1

	Experimental	Theoretical
F:C ratio	1.70 ± 0.3	1.75
% CF_2 group	$61\% \pm 2\%$	75%
% CF_3 group	$12\% \pm 2\%$	8%

The difference between theoretical and experimental CF_2 group and CF_3 group percentages can be attributed to a small amount of fragmentation of the perfluoroalkene monomer.

- 5 Figure 3 shows the C (1s) XPS spectrum for a 5 minute pulsed plasma polymerisation experiment where:- $P_{\text{cw}} = 50\text{W}$

$$T_{\text{on}} = 20\mu\text{s}$$

$$T_{\text{off}} = 10000\mu\text{s} \quad \langle P \rangle = 0.1\text{W}$$

10

The chemical composition of the deposited coating for pulsed plasma deposition is given in Table 2 below.

Table 2

15

	Experimental	Theoretical
F:C ratio	1.75 ± 0.7	1.75
% CF_2 group	$63\% \pm 2\%$	75%
% CF_3 group	$10\% \pm 2\%$	8%

- It can be seen that the CF_2 region is better resolved and has greater intensity which means less fragmentation of the perfluoroalkyl tail compared to continuous wave plasma
- 20 polymerisation.

- Surface energy measurements were carried out on slides produced in this way using dynamic contact angle analysis. The results showed that the surface energy was in the range
- 25 of $5\text{-}6\text{mJm}^{-2}$.

Example 2Oil and Water Repellency Test

The pulsed plasma deposition conditions described in Example 1 above were used to coat a piece of cotton (3x8cm) which was then tested for wettability using "3M Test Methods" (3M oil repellency Test 1, 3M Test Methods Oct.1, 1988). As a Water repellency test, the 3M water repellency Test II, water/alcohol drop test, 3M Test 1, 3M Test Methods, October 1, 1988 was used. These tests are designed to detect a fluorochemical finish on all types of fabrics by measuring:

(a) aqueous stain resistance using mixtures of water and isopropyl alcohol.

(b) the fabric's resistance to wetting by a selected series of hydrocarbon liquids of different surface tensions.

These tests are not intended to give an absolute measure of the fabric's resistance to staining by watery or oily materials, since other factors such as fabric construction, fibre type, dyes, other finishing agents, etc., also influence stain resistance. These tests can, however, be used to compare various finishes. The water repellency tests comprises placing 3 drops of a standard test liquid consisting of specified proportions of water and isopropyl alcohol by volume onto the plasma polymerised surface. The surface is considered to repel this liquid if after 10 seconds, 2 of the 3 drops do not wet the fabric. From this, the water repellency rating is taken as being the test liquid with the greater proportion of isopropyl alcohol which passes the test. In the case of the oil repellency test, 3 drops of hydrocarbon liquid are placed on the coated

surface. If after 30 seconds no penetration or wetting of the fabric at the liquid-fabric interface occurs around 2 of the 3 drops is evident, then the test is passed.

- 5 The oil repellency rating is taken to be the highest-numbered test liquid which does not wet the fabric surface (where the increasing number corresponds to decreasing hydrocarbon chain and surface tension).
- 10 The ratings obtained for the pulsed plasma deposition of 1H, 1H, 2H perfluoro-1-dodecene onto cellulose were:-

Water	9	(10% water, 90% isopropyl alcohol)
Oil	5	(dodecane)

15

These values compare well with commercial treatments.

Example 3

Plasma Polymerisation of Acrylates

- 20 The method of Example 1 described above was repeated using 1H, 1H, 2H, 2H-heptadecafluorodecyl acrylate (Fluorochem F04389E, 98% purity) in place of the perfluoroalkene. As in Example 1, low average powers were used for continuous wave and pulsed plasma polymerisation experiments. For example,
- 25 the XPS spectrum of a 1W continuous wave plasma polymer deposited onto a glass slide for 10 minutes is shown in Figure 4(a). Figure 4(b) shows the C(1s) XPS spectrum for a 10 minute pulsed plasma polymerisation experiment where
- 30 $P_{cw} = 40W$ (average continuous wave power)
 $T_{on} = 20\mu s$ (pulsed time on)
 $T_{off} = 20000\mu s$ (pulsed time off)
 $\langle P \rangle = 0.04W$ (average pulsed power)

17

Table 3 compares the theoretical (taken from the monomer, $\text{CH}_2=\text{CHCO}_2\text{CH}_2\text{CH}_2\text{C}_6\text{F}_{11}$) environments with what is actually found for polymer coatings.

5

Table 3

Environment	eV	Theoretical percentages	Experimental percentages
CF_3	293.2	7.7	7.8
CF_2	291.2	53.8	47.0
$\text{O}-\text{C}=\text{O}$	289.0	7.7	13.0
CF	287.8	--	0.7
$\text{C}-\text{CF}_n/\text{C}-\text{O}$	286.6	15.4	13.4
$\text{C}-\text{C}(\text{O})=\text{O}$	285.7	7.7	3.9
C_xH_y	285.0	7.7	7.2

It can be seen that the CF_2 group is the prominent environment in the C(1s) XPS envelope at 291.2eV. The remaining carbon environments being CF_3 , partially fluorinated and oxygenated carbon centres and a small amount of hydrocarbon (C_xH_y). The chemical composition of the coatings deposited for continuous wave and pulsed plasma conditions are given below in Table 4 (excluding satellite percentages) along with the theoretically expected compositions).

15

Table 4

	Theoretical	CW Plasma	Pulsed Plasma
F:C ratio	1.31	0.94	1.49
% CF_2 group	53.8%	27.2%	47.0%
% CF_3 group	7.7%	3.8%	7.8%

20

It can be seen from Figure 4(b) that the CF_2 region is better resolved and has greater intensity, which means less fragmentation of the perfluoroalkyl tail occurs during pulsed plasma conditions compared to continuous wave plasma polymerisation. In the case of the continuous wave plasma experiments, the low percentages of CF_2 and CF groups occur.

Surface energy measurements as described in Example 1 shows a surface energy of $6mJm^{-1}$.

Example 4

Oil and Water Repellency Test

Using the pulsed plasma deposition conditions of Example 3 except that these were applied for 15 minutes, pieces of cotton (3x 8cm) were coated with 1H, 1H, 2H, 2H-heptadecafluorodecyl acrylate. Similar pieces of cotton were coated with the same compound using a continuous wave at 1W for 15 minutes. These were then subjected to oil and water repellency tests as described in Example 2 above.

Samples were then subjected to a benzotrifluoride Soxhlet extraction for either 1 or 7 hours and the oil and water repellency tests repeated. The results, expressed as described in Example 2,

Time (hours)	Continuous wave		Pulsed wave	
	Oil- repellency	Water repellency	Oil repellency	Water repellency
0	7	4	8	10
1	-	2	6	7
7	-	2	5	7

Hence these coatings are highly hydrophobic and oleophobic and the coatings have good durability.

5

Example 5

Treatment of silicone coated synthetic fabric

A sample of a modified acrylic/nylon fabric which already contained a silicone coating to impart water repellency, was
10 subjected to the a pulsed acrylate plasma consisting of the compound $\text{CH}_2=\text{CHCOO}(\text{CH}_2)_4\text{C}_6\text{F}_{13}$, and using the conditions described in Example 3.

A sample of the same material was subjected to a two stage
15 deposition process in which the fabric was first exposed to a continuous wave 30W air plasma for 5 seconds followed by exposure to the same acrylate vapour only.
The products were then tested for oil and water repellency as described in Example 2.

20

In addition, the durability of the coating was tested by then subjecting the products to a 1 hour Soxhlet extraction with trichloroethylene.

25 The results are as shown in Table 5

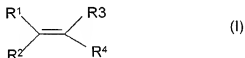
Table 5

Treatment	Repellency Ratings		
	Before Plasma	After Plasma	After extraction with solvent
Pulsed phase acrylate plasma	W2	O7, W10	O6, W8
Air plasma followed by exposure to acrylate monomer	W2	O1, W3	O1(borderline) W2

- 5 It appears therefore that the process of the invention can not only enhance the water repellency of such as fabric, and also confer oil repellency, the durability of the coating is higher than that obtained using the known two step grafting polymerisation process.

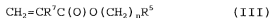
Claims

1. A method of coating a surface with a polymer layer, which method comprises exposing said surface to a plasma comprising a monomeric unsaturated organic compound which comprises an optionally substituted hydrocarbon group wherein any optional substituents are halogen; provided that where the compound is a straight chain perhalogenated alkene, it comprises at least 5 carbon atoms; so as to form an oil or water repellent coating on said substrate.
2. A method according to claim 1 wherein the organic compound is substituted by halogen.
3. A method according to claim 1 wherein the said organic compound is a compound of formula (I)



- where R^1 , R^2 and R^3 are independently selected from hydrogen, alkyl, haloalkyl or aryl optionally substituted by halo; and R^4 is a group $X-R^5$ where R^5 is an alkyl or haloalkyl group and X is a bond; a group of formula $-C(O)O(CH_2)_nY-$ where n is an integer of from 1 to 10 and Y is a bond or a sulphonamide group; or a group $-(O)_pR^6(O)_q(CH_2)_t-$ where R^6 is aryl optionally substituted by halo, p is 0 or 1, q is 0 or 1 and t is 0 or an integer of from 1 to 10, provided that where q is 1, t is other than 0.
4. A method according to claim 3 wherein R^5 is a haloalkyl group.

5. A method according to claim 4 wherein R^3 is a perhaloalkyl group.
6. A method according to claim 5 wherein R^5 is a perfluoroalkyl group of formula C_mF_{2m+1} where m is an integer of 1 or more.
7. A method according to claim 6 wherein m is from 1-20.
8. A method according to claim 7 wherein m is from 6-12.
9. A method according to any one of the preceding claims wherein R^1 , R^2 and R^3 are independently selected from hydrogen or a C_{1-6} alkyl or halo C_{1-6} alkyl group.
10. A method according to claim 9 wherein at least one of R^1 , R^2 and R^3 is hydrogen.
11. A method according to claim 10 wherein R^1 , R^2 , R^3 are all hydrogen.
12. A method according to claim 3 wherein X is a group of formula $-C(O)O(CH_2)_nY-$ and Y is a sulphonamide group of formula $-N(R^6)SO_2-$ where R^6 is hydrogen or alkyl.
13. A method according to claim 3 wherein the compound of formula (I) comprises a compound of formula (II)
- $$CH_2=CH-R^5 \quad (II)$$
- where R^5 is as defined in claim 3.
14. A method according to claim 3 wherein the compound of formula (I) is an acrylate of formula (III)



where n and R⁵ as defined in claim 2 and R⁷ is hydrogen or
5 C₁₋₆ alkyl.

15. A method according to any one of the preceding claims
wherein the surface is a surface of a fabric, metal, glass,
ceramics, paper or polymer substrate.

10

16. A method according to claim 15 wherein the substrate
is a fabric.

17. A method according to any one of the preceding claims
15 wherein the gas pressure of the organic compound is from
0.01 to 10 mbar.

18. A method according to any one of the preceding claims
wherein a glow discharge is ignited by applying a high
20 frequency voltage.

19. A method according to claim 18 wherein the voltage is
applied as a continuous field.

25 20. A method according to claim 18 wherein the voltage is
applied as a pulsed field.

21. A method according to claim 20 wherein pulses are
applied in a sequence which yields low average power.

30

22. A method according to claim 21 wherein the sequence is
such that the power is on for 20µs and off for from 10000µs
to 20000µs.

23. A method according to any one of the preceding claims wherein the plasma polymerisation takes place for from 2 to 15 minutes.
- 5 24. A method according to claim 1, which method comprises exposing a surface to a plasma comprising a compound of formula (III) as defined in claim 14, said plasma being created by a pulsed voltage.
- 10 25. A hydrophobic or oleophobic substrate which comprises a substrate comprising a coating of a polymer which has been applied by the method according to any one of claims 1 to 24.
- 15 26. A hydrophobic or oleophobic substrate according to claim 25 wherein the polymer is a haloalkyl polymer.
27. A substrate according to claim 25 or claim 26 which is a fabric.
- 20 28. An item of clothing which comprises a fabric according to claim 27.

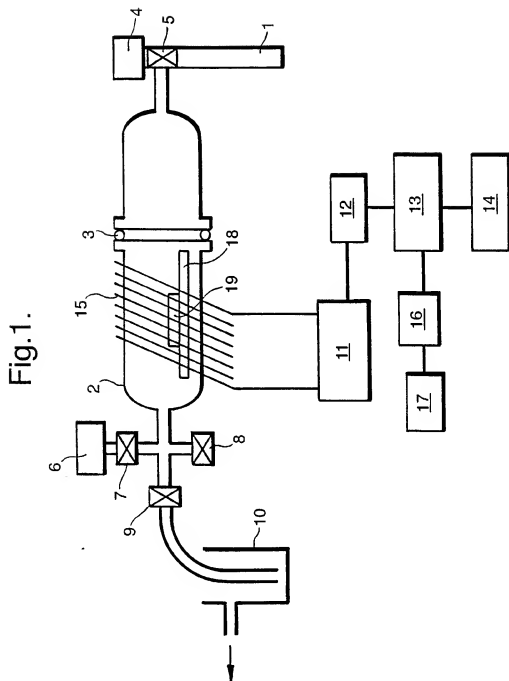


Fig.2.

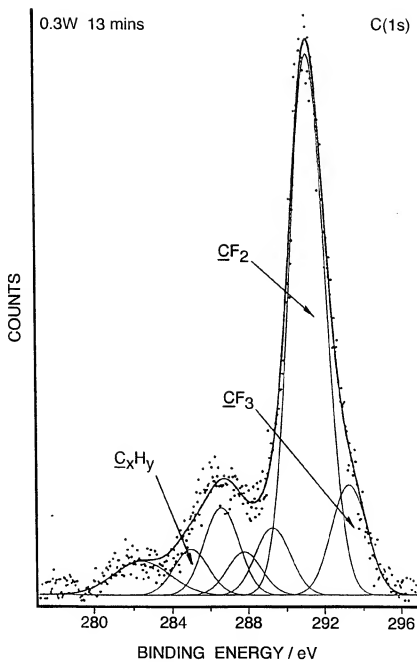


Fig.3.

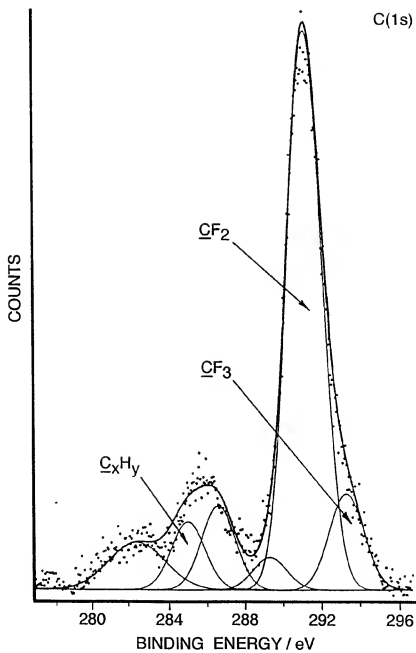
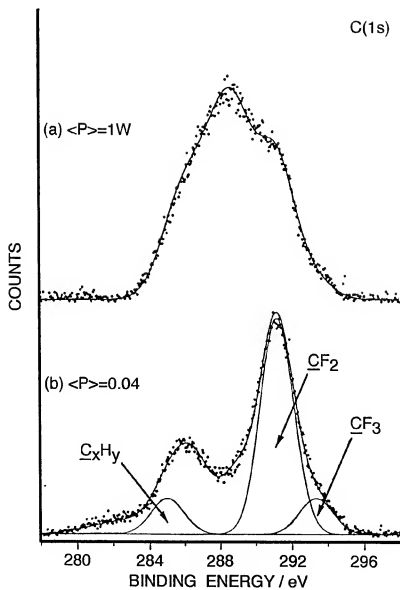


Fig.4.



INTERNATIONAL SEARCH REPORT

International Application No.

PCT/GB 98/01702

A. CLASSIFICATION OF SUBJECT MATTER

IPC 6 D06M10/02 D06M14/18 D21H19/16 D21H23/44 B29C59/14

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 6 D06M D21H B29C B05D

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	<p>DATABASE WPI Week 8235 Derwent Publications Ltd., London, GB; AN 82-73505E XP002079483 & JP 57 119906 A (DAIKIN KOGYO KK) , 26 July 1982 see abstract</p> <p style="text-align: center;">---</p>	<p>1-11, 14, 15, 17, 18, 25, 26</p>
X	<p>PATENT ABSTRACTS OF JAPAN vol. 014, no. 242 (C-0721), 23 May 1990 & JP 02 061176 A (WAKAYAMA PREFE CTURE GOV), 1 March 1990 see abstract</p> <p style="text-align: center;">---</p> <p style="text-align: center;">-/--</p>	<p>1, 15, 16, 25, 27, 28</p>

☒

Further documents are listed in the continuation of box C.

☒

Patent family members are listed in annex.

* Special categories of cited documents:

"A" document defining the general state of the art which is not considered to be of particular relevance

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"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

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Date of the actual completion of the international search

5 October 1998

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INTERNATIONAL SEARCH REPORT

Int. l. Application No.

PCT/GB 98/01702

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	EP 0 049 884 A (DAIKIN IND LTD) 21 April 1982 cited in the application see page 1, line 22 - page 4, line 11 see page 5, line 16 - page 6, line 32 ---	1-11,14, 15,17-26
X	US 5 041 304 A (KUSANO YUKIHIRO ET AL) 20 August 1991 see column 2, line 22 - column 5, line 29; claims ---	1,2,15, 16,25-28
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X	US 5 290 378 A (KUSANO YUKIHIRO ET AL) 1 March 1994 see column 2, line 47 - column 5, line 10; claims -----	1,15

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International Application No

PCT/GB 98/01702

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